UP AND UP ALLERGY RELIEF- loratadine tablet
Target Corporation

Target Corporation Allergy Relief Drug Facts

Active ingredient (in each tablet)

Loratadine 10 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- itchy, watery eyes
- sneezing
- itching of the nose or throat

Warnings

Do not use

if you have ever had an allergic reaction to this product or any of its ingredients

Ask a doctor before use if you have

liver or kidney disease. Your doctor should determine if you need a different dose.

When using this product

do not take more than directed. Taking more than directed may cause drowsiness.

Stop use and ask a doctor if

an allergic reaction to this product occurs. Seek medical help right away.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

Directions

adults and children 6 years and over	1 tablet daily; not more than 1 tablet in 24 hours
children under 6 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

Other information

- do not use if printed foil under cap is broken or missing
- store between 20° to 25°C (68° to 77°F)

Inactive ingredients

lactose monohydrate, magnesium stearate, povidone, pregelatinized starch

Questions?

Call 1-888-547-7400

Principal Display Panel

CE PACK

Compare to active ingredient in Claritin®

non-drowsy*

allergy relief

loratadine tablets, 10 mg

antihistamine

30 days of relief

original prescription strength

indoor and outdoor allergies

24 hour relief of:

sneezing

runny nose

itchy, watery eyes

itchy throat or nose

24 HOUR

ACTUAL SIZE

30 TABLETS

30 TABLETS

*When taken as directed. See drug facts panel.

UP AND UP ALLERGY RELIEF

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-612
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	LORATADINE	10 mg

Inactive Ingredients

Ingredient Name	Strength
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POVIDONE (UNII: FZ989GH94E)	

Product Characteristics

Color	WHITE	**Score**	no score
Shape	OVAL	**Size**	8mm
Flavor		**Imprint Code**	L612
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-612-46	10 in 1 CARTON	03/26/2012	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:11673-612-65	1 in 1 CARTON	04/12/2012	
2		30 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:11673-612-72	1 in 1 CARTON	04/16/2012	07/01/2016
3		60 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:11673-612-76	1 in 1 CARTON	04/12/2012	02/01/2016
4		120 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:11673-612-82	1 in 1 CARTON	04/12/2012	
5		200 in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:11673-612-95	1 in 1 CARTON	02/27/2013	02/27/2013
6		45 in 1 BOTTLE; Type 0: Not a Combination Product		
7	NDC:11673-612-03	1 in 1 CARTON	02/19/2015	
7		70 in 1 BOTTLE; Type 0: Not a Combination Product		
8	NDC:11673-612-78	100 in 1 BOTTLE; Type 0: Not a Combination Product	03/20/2015	
9	NDC:11673-612-58	1 in 1 CARTON	03/25/2015	
9		40 in 1 BOTTLE; Type 0: Not a Combination Product		
10	NDC:11673-612-87	1 in 1 CARTON	03/15/2016	
10		300 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076301	03/26/2012	

Labeler - Target Corporation (006961700)